

Invited Review

The effectiveness of digital interventions for self-management of chronic pain in employment settings: a systematic review

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Introduction: Chronic pain affects over a quarter of the workforce with high economic burden for individuals, employers and healthcare services. Access to work-related advice for people with chronic pain is variable. This systematic review aims to explore the effectiveness of workplace-delivered digital interventions for the self-management of chronic pain.

Source of data: MEDLINE, EMBASE, CINAHL, PsycINFO, the Cochrane Library, JBI, Open Science Framework, Epistemonikos and Google Scholar. Articles published between January 2001 and December 2023 were included. Searches were conducted between October 2023 and December 2023.

Areas of agreement: Workplace-delivered digital interventions to support self-management of chronic pain at work may improve pain and health-related quality of life in vocationally active adults. Delivering interventions outside of clinical services, through the workplace setting, may help to reduce inequity in access to work-related advice for people with chronic pain, and ultimately reduce the burden on individuals, employers and healthcare services. Interventions include mobile apps and web-based programmes.

Areas of controversy: Studies were moderate-to-low quality. Most studies focused on exercise, few considered other aspects of pain self-management.

Given the limited evidence in the current literature, consensus on best intervention format and delivery is lacking.

Growing points: More high-quality studies are needed given the heterogeneity in study design, interventions and outcome measures.

Areas timely for developing research: No interventions included advice on work-related adjustments or support. Few studies included work-related outcomes, despite the known impact of pain on work and work on health.

Key words: chronic pain, workplace, self-management, digital, occupational health, health promotion

Introduction

Chronic pain is a global health priority. Prevalence estimates across 52 countries range from 9.9% to 50.3%¹ with a high economic burden for individuals, employers and healthcare services (over £100 billion per annum in the UK²). Chronic pain can impact on people's ability to work, their productivity, sickness absence, presenteeism and early retirement due to disability^{3,4}. Retaining people with chronic pain in the workforce is important since unemployment is associated with an increased risk of mortality and morbidity, and good work improves health and wellbeing and reduces social exclusion.⁵ Providing advice and information to people with chronic pain about self-management strategies is recommended within clinical guidelines for chronic pain management.⁶ Self-management is equipping patients 'with skills to actively participate and take responsibility in the management of their chronic condition in order to function optimally' and may involve a combination of knowledge acquisition, sign/symptom monitoring, medication management, enhancing problem-solving/decision-making skills for medical treatment management and/or changing health behaviour(s).⁷ Self-management advice is routinely provided by healthcare professionals, but this rarely includes discussion about self-management strategies in the context of work. Although work-related self-management is a core focus of occupational therapy (OT), access to OT services is highly variable,⁸ meaning that many

people with chronic pain do not receive work-related self-management advice. One route to supporting people to managing chronic pain at work (and potentially reducing burden on healthcare services) is to offer self-management interventions through employment settings. In the UK, around three-quarters of people aged 16–64 years are in employment. Given the high prevalence of chronic pain (one-third to one-half of the population), workplace-delivered interventions have potential for wide reach. Additionally, targeting interventions through non-clinical settings, such as workplaces, could help to reduce inequity in access to work-related advice and support through healthcare services. Digital interventions (DIs) are potentially scalable⁹ and may facilitate in reaching those with chronic pain regardless of their activity level, pain status, occupation type or geographical location. DIs provide information and/or support (emotional, decisional and/or behavioural) via digital platforms (e.g. website, computer, tablet or smartphone). Although workplace-focused DIs are emerging (e.g. Blake *et al.*¹⁰), the effectiveness of workplace delivered DIs in reducing pain, improving health, wellbeing, quality of life and work-related outcomes has not yet been established.

Study aim

To conduct a systematic literature review to explore the effectiveness of DIs for self-management of chronic pain in employment settings.

Methods

This systematic review was pre-registered with PROSPERO on October 19, 2023 (CRD42023463484).

Eligibility criteria

All original studies consisting of randomized control trials (RCTs) and repeated measures non-randomized trials (RMs). The trials all included a DI and were conducted with vocationally active adult participants. The DI should function without any direct input from health professionals and require interaction with the participant. All studies were published since 2001, the year the term electronic health (eHealth) first emerged.¹¹ Participants were recruited via their workplace. Articles were restricted to the English language, but there were no geographical limitations. Studies were excluded where the intervention was solely an appointment reminder or treatment compliance or telehealth or via email or direct input with a practitioner. Studies involving only passive monitoring (e.g. step counters only) or only reminders were excluded. Reviews, opinions, letters and unpublished literature were not considered.

Search strategy

This systematic review was conducted according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses: the 2020 PRISMA statement¹² (Fig. 1). The following databases were searched electronically: MEDLINE, EMBASE, CINAHL, PsycINFO, the Cochrane Library, JBI, Open Science Framework, Epistemonikos and Google Scholar. Searches were conducted between October 2023 and December 2023. The search strategy (Supplementary File S1) was developed with a combination of Medical Subject Headings and keywords and using filters from the Cochrane Back Review Group. References of selected articles were hand-searched for eligible studies. A search of Open Grey and Google Scholar revealed materials with reference lists relevant to this review.

Study selection

Two authors (W.J.C., H.B.) were involved in study selection. Records were managed through Covidence systematic review software (Veritas Health Innovation, Melbourne, Australia). An initial screening of titles and abstracts of studies retrieved was conducted (W.J.C.), to identify studies that meet the study inclusion criteria outlined above. A second reviewer (H.B.) independently screened 20% of titles and abstracts. Full text was obtained for abstracts with insufficient information or in a situation of disagreement. A study was included when both reviewers independently assessed it as satisfying the inclusion criteria from the full text. Any disagreements were resolved through discussion.

Data items

Three authors (W.J.C., A.G. and H.B.) were involved in data extraction. Data extraction was independently performed on all included articles by two authors (W.J.C. and A.G.), a 20% check was conducted by a third author (H.B.). The following data were extracted: author and year, name of the journal, study design, inclusion/exclusion criteria, number of participants, participant characteristics (age, gender ratio), pain location, type of intervention, intervention duration, outcome measures (pain, quality of life, psychological, behavioural, physical activity, employment measures, other).

Study risk of bias assessment

Risk of bias for each included trial was independently assessed by two reviewers (W.J.C. and A.G.) using the revised JBI critical appraisal tool for randomized controlled trials and the equivalent for quasi-experimental studies or non-randomized trials.¹³

Strategy for data synthesis

We provide a narrative synthesis of the findings from the included studies, structured around study designs and settings, target population characteristics, type of intervention, intervention content and type of

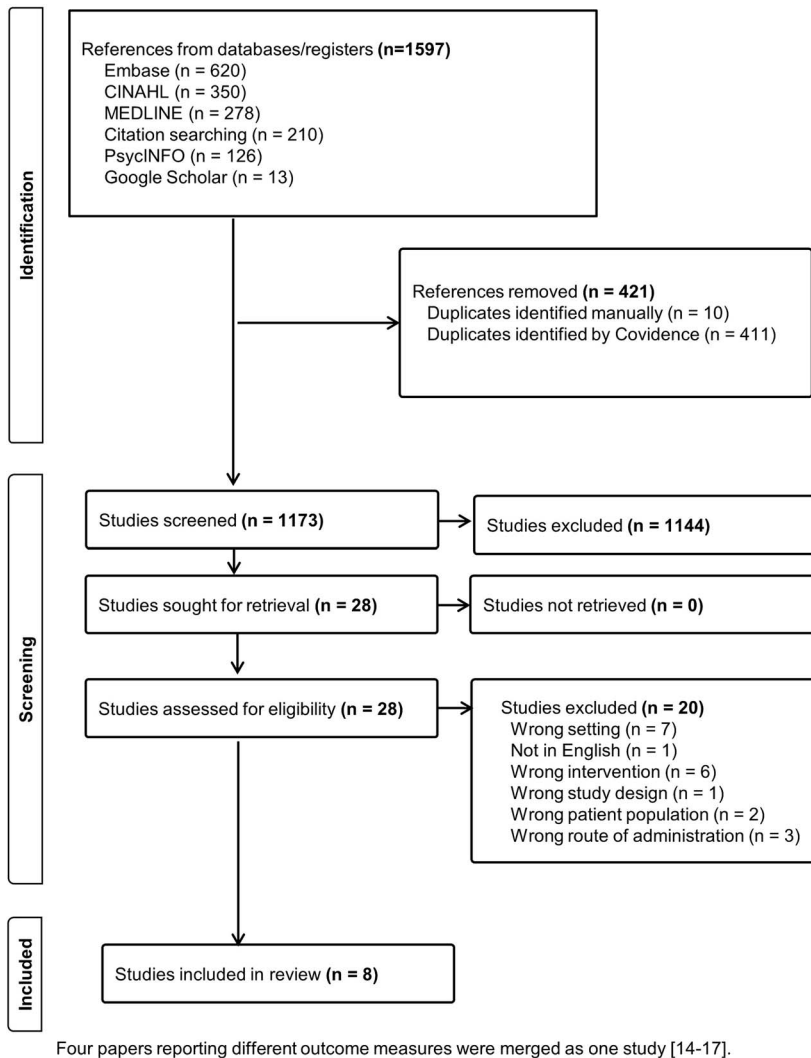


Fig. 1 Flow chart showing article selection strategy, including reasons for exclusion according to the PRISMA guidelines.

outcome. We provide summaries of intervention effects for each study by reporting between-group differences.

Results

Study selection

The search identified 1591 records and 28 full articles that met the inclusion criteria for further examination. Twenty articles were excluded. The findings of four articles were merged as they reported on

a single study,¹⁴⁻¹⁷ and the remaining seven articles were selected for inclusion in the review. The articles were published between 2011 and 2021. The flowchart of the literature search is shown in [Figure 1](#). Participant demographics and summary of interventions are shown in [Table 1](#).

Study designs and settings

The eight intervention studies¹⁴⁻²¹ included five RCTs, of which two were pilot RCTs^{18,21} and one was a cluster RCT,²⁰ and three repeated measures (RMs)

Table 1 Participant demographics and intervention type

Study # with ref.	Design	Partici pants n	Age, mean (SD)	Males n (%)	Digital intervention	Duration	Exercise/ stretch	Breaks	Mind-fulness	Posture	Other
1 ¹⁸	RCT-pilot	41	C:41.7 (6.35), I: 40.48 (7.22)	C: 12 (29), I:7 (17.1)	Smartphone	6 weeks	✓	✓			
2 ¹⁹	RCT	121	C: 42.4 (8.7), I: 42.4 (8.0)	C:33 (72), I: 39 (81)	Smartphone AI-assisted chatbot	12 weeks	✓		✓	✓	Exercise motivation nudges
3 ¹⁴⁻¹⁷	RCT	100	C: 45.5 (7.02), I: 46.83 (9.13)	C:11 (4%), I:15.2%	Web-based	9 months	✓			✓	
4 ²⁰	Cluster RCT	175	C:42.9 (12.0), I:41.8 (11.4)	C:77 (97.5), I: 85 (88.5)	Computer-based	4 months	✓	✓			Education worksheets
5 ²¹	RCT-pilot	21	C:27.56 (4.67), I: 27.09 (4.83)	C:6 (55) I: 5 (45)	Smartphone	8 weeks	✓				
6 ²²	Repeated measures	30	28.13 (2.97)	13 (66.50)	Smartphone	8 weeks	✓				
7 ²³	Repeated measures	645	56 (12.83)	198 (30.7)	Web-based	6 months					Setting expectations, managing stress, coping with pain, accessing social support, healthy sleep, nutrition, exercise, improving doctor-patient relationships, medication adherence and chronic disease self-management.
8 ²⁴	Repeated measures	417	49.8	128(30.7)	Web-based	30, 90, 180 days					Digital health coaching sources of stress, perceived barriers to managing stress, coping skills and resources and stages of change.

C, control; I, intervention web-based applications are accessed via the internet, computer-based applications were a software programme loaded onto an individual computer.

Table 2 Outcome variables

Study # lead author	Pain location	Inclusion	Exclusion	Pain	Quality of life	Psychological	Physical Activity	Behavioural	Employment metrics	Other
1 Alimhdawi ¹⁸	LBP	Aged 30–55 years, office worker >5y > 5 h daily desk work.	Pregnancy, specific back diagnosis	VAS B: 5.62 ± 2.06 FU: 2.30 ± 2.13, P < 0.001 ODI B: 30.95 ± 9.31 FU: 20.25 ± 13.47, P = 0.002 Cohen's d = 1.71	SF-12: PCS, MCS B: 67.67 ± 17.64 F: 79.95 ± 16.09 P = 0.001 Cohen's d = 0.18 PCS Cohen's d = 1.08 MCS Cohen's d = 0.131	DASS Cohen's d Stress = 0.34, Cohen's d Anxiety = 0.18 Cohen's d Depression = 0.20	IPAQ Cohen's d = 0.45	Hours, days, sickness, mean computer time, phone usage,	PSQI Cohen's d = 0.14	
2 Anan ¹⁹	Neck, shoulder and back	Screening questions and back smartphone	Pregnancy, CVD, other clinical trials, disability/exercise restriction	Subjective pain severity neck/shoulder and back + subjective pain stiffness neck/shoulder and back + improvement 36 (76%) improved P < 0.001						Adherence rate 92% (44/48)
3 del Pozo-Cruz ^{14–17} (four articles, 1 study)	Back	18–64 years, + diagnosis of subacute LBP physical activity < 2 working on a computer daily	Diagnosed backache, reported chronic backache, clinical red flags for disc disease, any other major illness, lack of fluency in Spanish	ODI, OR 5.40 (95% CI 1.71, 17.22), P = 0.001 Cohen's d = 0.93 RDM B: 12.28 ± 2.63 F: 4.93 ± 2.59, P < 0.001 Cohen's d = 2.82 SBST (TOTAL) B: 4.38 (1.48) F: 3.39 (1.39), P = 0.019 Cohen's d = 0.98	EQ-5D-3L OR 3.59 (95% CI 2.21, 5.82), P < 0.001 Cohen's d = 0.71	SBST psych Fear avoidance. OR 0.35 (95% CI 0.15, 0.84), P = 0.017 Cohen's d = 0.58	Lumbar endurance B: 78.80 ± 30.6 F: 92.4 ± 27.9 P < 0.001 Cohen's d = 0.46 Abdominal endurance B: 48.1 ± 33.0 F: 64.4 ± 30.7, P < 0.001 Cohen's d = 0.51	Stage of change questionnaire		

(Continued)

Table 2 Continued

Study # lead author	Pain location	Inclusion	Exclusion	Pain	Quality of life	Psychologi- cal	Physical Activity	Behavioural	Employ- ment metrics	Other
4 Lanthers 20	Any	18–65 years + > 5 h VDU per day + no sickness absence in 1 month + admin staff no behavioural or learning disorder + no new VDU equipment	Maternity leave, change in job	Nordic questionnaire Reduction in 30 days, $P = 0.038$		HADS	IPAQ sedentary time per day		Stress at work, satisfaction at work,	DI acceptability, eye strain, corrective spectacles or lenses, exercise adherence
5 Lee 21	Neck	2.5–35 years 6 + computer hours neck pain for more than 6 months	<3 VAS pain A history of traumatic injury on neck, a congenital deformity, a history of surgical operation or injection on neck and any neurological symptoms.	VAS, B:5.20 ± 2.19 F:2.73 ± 1.99, $P < 0.05$ pain duration, McKenzie classification, Cohen's $d = 1.18$ Functional disability—neck disability index questionnaire, strength and ROM B: 26.80 ± 9.68 F:17.25 ± 8.34, $P < 0.05$ Cohen's $d = 1.06$	SF-36 PCS, B:43.18 ± 8.58 F: 48.40 ± 7.22, $P < 0.05$ Cohen's $d = 0.67$	Fear- avoidance (Work) B: 25.18 ± 3.97 F:20.73 ± 6.4 $P < 0.05$ Cohen's $d = 0.84$			Maximal voluntary strength Neck extension B:16.82 ± 7.74 F:25.92 ± 6.86, $P < 0.05$ Cohen's $d = 1.24$	

(Continued)

Table 2 Continued

Study # lead author	Pain location	Inclusion	Exclusion	Pain	Quality of life	Psychological	Physical Activity	Behavioural	Employment metrics	Other
6 Lee ²²	Neck	Neck pain > 3 months; mean pain > 3 in the last week; smartphone	(1) They had received any other treatment or surgery within 3 months; or (2) Their neck pain was caused by a known trauma, rheumatic disorder or malignant disease.	VAS, B:4.63 ± 1.89, F:2.0 ± 1.69, P < 0.001 Cohen's d = 1.47 McKenzie classification, Functional disability—Neck disability index questionnaire, strength and ROM B:22.18 ± 9.42 F:13.74 ± 7.28, P < 0.001 Cohen's d = 1.00.	SF-36 PCS, B: 46.14 ± 7.29, F:51.22 ± 6.55, P = 0.02 Cohen's d = 0.73	Fear-avoidance. (Work)	Exercise minutes per day and sedentary minutes per day		Hours, days, mean computer time, 3.91 ± 0.51/5	The patient satisfaction was 91.85% (37.78%), and the mean duration per exercise session was 16.8 ± 7.38 min
7 Nevedal ²³	Any	Eligible participants were either employed by 1/37 participating US companies or a member of 1 of 18 participating US healthcare plans.	This program isn't intended for people suffering from the following: acute pain, cancer pain, pelvic or abdominal pain.	VAS, B:5.30 ± 2.46, F:3.72 ± 2.73, P < 0.001 Cohen's d = 0.61 duration, McKenzie classification, Functional disability, neck pain, pain interference, pain unpleasantness, pain medication B:5.43 ± 2.52 F:3.78 ± 2.79, P < 0.001 Cohen's d = 0.62	Quality of life—CDC HRQOL-4 (1 item) B: <Fair 20.6% F: <Fair 16.5%, P = 0.006	Participant stress		Motivation to manage and confidence to manage pain	Quality of health—CDC HRQOL-4 (1 item)	

(Continued)

Table 2 Continued

Study #	Pain location	Inclusion	Exclusion	Pain	Quality of life	Psychological	Physical Activity	Behavioural	Employment metrics	Other
8	Any	First, subjects must have participated in: HealthMediaR CareTM for pain (a chronic pain management program);		Pain improvement and pain worsening were both associated with productivity impairment. $P < 0.001$		CES-D Boston Form Depression/10			Work productivity and activity impairment questionnaire B:37.62 F: 29.29, $P < 0.001$ Cohen's d unavailable	

Only significant findings are reported. **BOLD** items showed statistically significant change.

VAS, Visual Analogue Scale; ODI, Oswestry Disability Index; RDM, Roland-Morris Disability Questionnaire; SBST, StarT Back Tool; CES-D, Center for Epidemiologic Studies Depression Scale; SF-36, Short Form Survey; SF-12, Short Form Survey; PCS, Physical Component Score (SF-12); DASS, Depression Anxiety Stress Scale; ROM, Range of Movement; HADS, Hospital Anxiety and Depression Score; PSQI, Pittsburgh Sleep Quality Index; VDU, visual display unit; IPAQ International Physical Activity Questionnaire; Quality of life/health-CDC HRQL-4, Centers for Disease Control and Prevention Health-related Quality of Life 4 item Measure; B, baseline; F, follow-up; OR, odds ratios.

